Amendments to the Claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (currently amended) A method for the treatment or prophylaxis of a human infected with hepatitis B virus comprising administering in combination or alternation an effective amount of:

β-2-hydroxymethyl-5-(5-fluorocytosin-1-yl)-1,3-oxathiolane (β-L-FTC); 1-(2'-deoxy-2'-fluoro-β-L-arabinofuranosyl)-thymine (L-FMAU); and interferon;

or their pharmaceutically acceptable salts or prodrugs, independently optionally in pharmaceutically acceptable carriers.

- 2. (original) The method of claim 1, wherein the β-L-FTC is in substantially pure form.
- 3. (original) The method of claim 1, wherein the β -L-FTC is at least 90% by weight of the β -L-isomer.
- 4. (original) The method of claim 1, wherein the β-L-FTC is at least 95% by weight of the β-L-isomer.
- 5. (currently amended) The method of claim 1, wherein the interferon is selected from the group consisting of interferon alpha, pegylated interferon alpha, interferon alpha-2a, interferon alpha-2b, pegylated interferon alpha-2b, ROFERON®-A (interferon

alpha-2a), PEGASYS® (pegylated interferon alpha-2a), INTRON®A (Interferon alpha-2b), PEG-INTRON® (pegylated Interferon alpha-2b), interferon beta, interferon gamma, interferon tau, interferon omega, consensus interferon, INFERGEN [[(]]interferon alphacon-1[[)]], OMNIFERON [[(]]natural interferon[[)]], REBIF [[(]]interferon beta-la[[)]], omega interferon, oral interferon alpha, interferon gamma-lb, SUPERFERON [[(]]natural human multi-subtype IFN-alpha[[)]], and HUFERON [[(]]human IFN-beta[[)]].

- 6. (original) The method of claim 5, wherein the interferon is interferon alpha.
- 7. (original) The method of claim 5, wherein the interferon is interferon gamma.
- 8. (original) The method of claim 5, wherein the interferon is interferon beta.